

Assessment of ECG and Echo Findings among With Chronic Kidney Disease Patients

Dr. B. Suresh Reddy

Assistant Professor, Department of General Medicine, Maheshwara Medical college and Hospital, Chitkul, Isnapur, Patancheru, Sangareddy, Telangana, India.

*Corresponding author

Dr. B. Suresh Reddy

Article History

Received: 29.08.2017

Accepted: 04.09.2017

Published: 30.09.2017



Abstract: Cardiovascular abnormalities are the leading cause of morbidity and mortality in Chronic Kidney Disease (CKD) patients which includes left ventricular hypertrophy (LVH), left ventricular dilation and left ventricular systolic and diastolic dysfunction. Cardiovascular complications lead in all causes of mortality among patients with CKD, accounting for approximately 50% of deaths. The present study designed to identify Electrocardiographic and Echocardiographic changes in patients with chronic kidney disease. A total of 50 CKD patients (23 males & 17 females) attending outpatient department were included. All patients were undergone for complete haemogram, ECG, Abdominal USG and Echocardiography. The results of present study showing, LVH with or without strain pattern in 62.5%, low voltage complexes in 15%, occasional ventricular premature complexes in 22.5% and non-specific ST – T changes in 30% of cases. 22.5% of Patients showed dilated LVH and 47.5% of showed concentric LVH. Systolic dysfunction was found in 27.5% of cases and diastolic dysfunction was found in 50% of cases. LVH is the commonest morphological abnormality observed. Echocardiographic study plays an important role in evaluating cardiac structure and functions as well as in stratifying prognostic risk. Periodic echocardiographic examination for diagnosis and treatment of cardiac abnormalities in patients with CKD is highly recommended.

Keywords: Chronic kidney disease (CKD), Echocardiography, Left ventricular hypertrophy (LVH), Systolic dysfunction, Diastolic dysfunction

INTRODUCTION

Chronic kidney disease (CKD), ranges from asymptomatic to total kidney failure is being widely alarming in India. CKD is characterized by reduced estimated glomerular filtration rate (eGFR) <60ml/min/1.73m² for more than 3 months and by structural or functional abnormalities [1, 2]. Cardiovascular complications are commonly encountered symptoms in CKD or end stage renal disease (ESRD) patients including left ventricular hypertrophy (LVH), Systolic and diastolic dysfunction. In addition, patients with CKD have a high prevalence of traditional and non-traditional risk factors such as diabetes mellitus, hypertension, dyslipidemia Uremia, inflammation and oxidative stress [3].

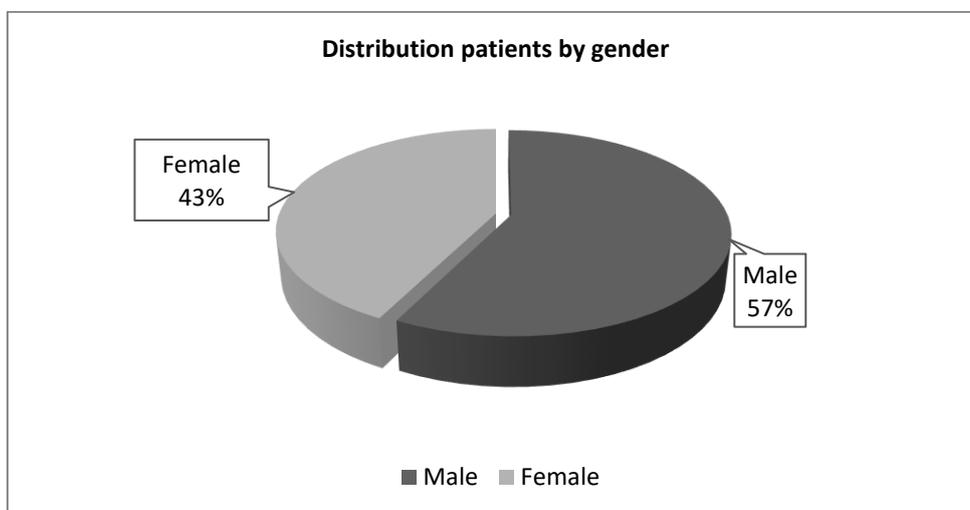
With the advancement of technology, Electrocardiogram (ECG) and Echocardiograph remains an essential tool for evaluation of cardiovascular disease [4, 5]. Echocardiographic study may play a critical role in evaluating cardiac functions especially for the assessment of left ventricular and right ventricular function [6].

From the above facts it is evident that, often there will be misinterpretation between CKD and CVD on clinical examination. It needs to be re-evaluated by easily available diagnostic procedures to prevent mortality and morbidity. The present was designed to investigate the electrocardiographic and echocardiographic changes in chronic kidney disease patients.

MATERIALS AND METHODS

The present study was conducted in Department of General medicine, Maheshwara Medical College and Hospital, Patancheru during June 2016 to April 2017. A total 50 patients (23 male & 17 female) with chronic kidney disease attending outpatient department were considered. Patients with GFR of 30-59 ml/min and with CKD were included and with Valvular Heart disease, Coronary artery disease, Systemic Hypertension on regular treatment and with poor pulmonary function was excluded from the study. All patients were undergone for complete haemogram, ECG, Abdominal USG and Echocardiography.

RESULTS



Graph-1: Distribution patients by sex

Table-1: Distribution of cases according to age group

Age (in years)	Males		Females		Total	
	No.	Percentage	No.	Percentage	No.	Percentage
0-20	-	-	1	2.5%	1	2.5%
21-30	-	-	-	-	-	-
31-40	3	7.5%	3	7.5%	6	15%
41-50	8	20%	6	15%	14	35%
51-60	6	15%	6	15%	12	30%
> 60	6	15%	1	2.5%	7	17.5%

Based on etiological distribution of patients 36% with diabetes mellitus, 14% with chronic glomerulonephritis, 12% chronic interstitial nephritis, 8%

with obstructive nephropathy, 6% with hypertension and 4% with autosomal dominant polycystic kidney diseases. Diabetes Mellitus is the commonest etiology.

Table-2: EEG changes in chronic kidney disease patients.

S.N	Finding	No. Of patients	Mean
1.	LVH	25	62.5%
2.	LVC	6	15%
3.	ST-T Changes	12	30%
4.	LBBB	1	2.5%
5.	VPC	9	22.5%
6.	Ischemia	24	60%

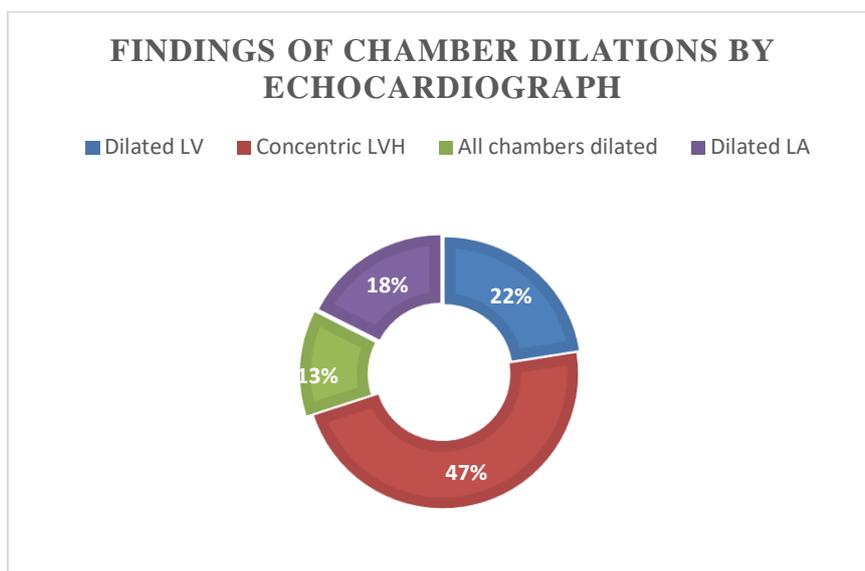


Fig-2: Echocardiographic finding in chronic kidney disease patients.

Table-3: Grading of regurgitations and LV function in CKD patients.

Grading	Mitral Regurgitation	Aortic Regurgitation	Tricuspid Regurgitation
Trivial	4	3	1
Mild	7	4	4
Moderate	1	-	-

Systolic dysfunction was found in 27.5% of cases, Type 1 diastolic dysfunction was found in 50% of cases.

DISCUSSION

Kidney and Heart are inseparably linked in terms of hemodynamic and regulatory functions. Their association exists at multiple levels including the renin-angiotensin aldosterone system, antidiuretic hormone, endothelin, the sympathetic nervous system and the natriuretic peptides. Globally, 40% of deaths in dialysis cases occurring by CVD [7].

In the present study, electrocardiogram showed evidence of LVH with or without strain pattern in 62.5%, low voltage complexes in 15%, occasional ventricular premature complexes in 22.5% and non-specific ST – T changes in 30% of cases. 22.5% of Patients showed dilated LVH and 47.5% of showed concentric LVH. The patients who have had long standing H/O Hypertension showed concentric LVH. These findings are comparable with studies done by NP Singh et al (76.92%), Foley *et al.* (73.9%) and Harnett *et al.* [8-10]. The prevalence of LVH in present study was observed in 62.5% of total cases. Study by McGregor et al found prevalence of LVH in 64-70%

of males and 63-65% of females and by P. Danger *et al* found higher prevalence in females. Studies suggest that LVH is the strongest independent predictable factor of adverse cardiovascular signs [11-14]. In present study non-specific ST – T changes in 30% of cases which is comparable with Shakira OM *et al* (22-39%) [15].

In the present study, aortosclerosis and posterior mitral annulus calcification was found in 35% of cases. Age, duration and hyperparathyroidism have been cited as prime determinants of valvar calcification [16, 17]. Systolic dysfunction was seen in 27.5% of cases, Diastolic dysfunction was seen in 50% of cases. LV diastolic dysfunction was more common in all stages of CKD [18]. According to Kramer *et al*, factors contributing to development of CCF in patient with CKD are Volume over load, Valvar heart disease, Negative inotropic effects of Calcium, Cardiac arrhythmias, Pressure overload, Myocardial damage, Anemia [19]. In present study, patients with moderate LV dysfunction showed features of volume overload, anemia and long standing history. All our patients revealed type I relaxation abnormality of diastolic dysfunction.

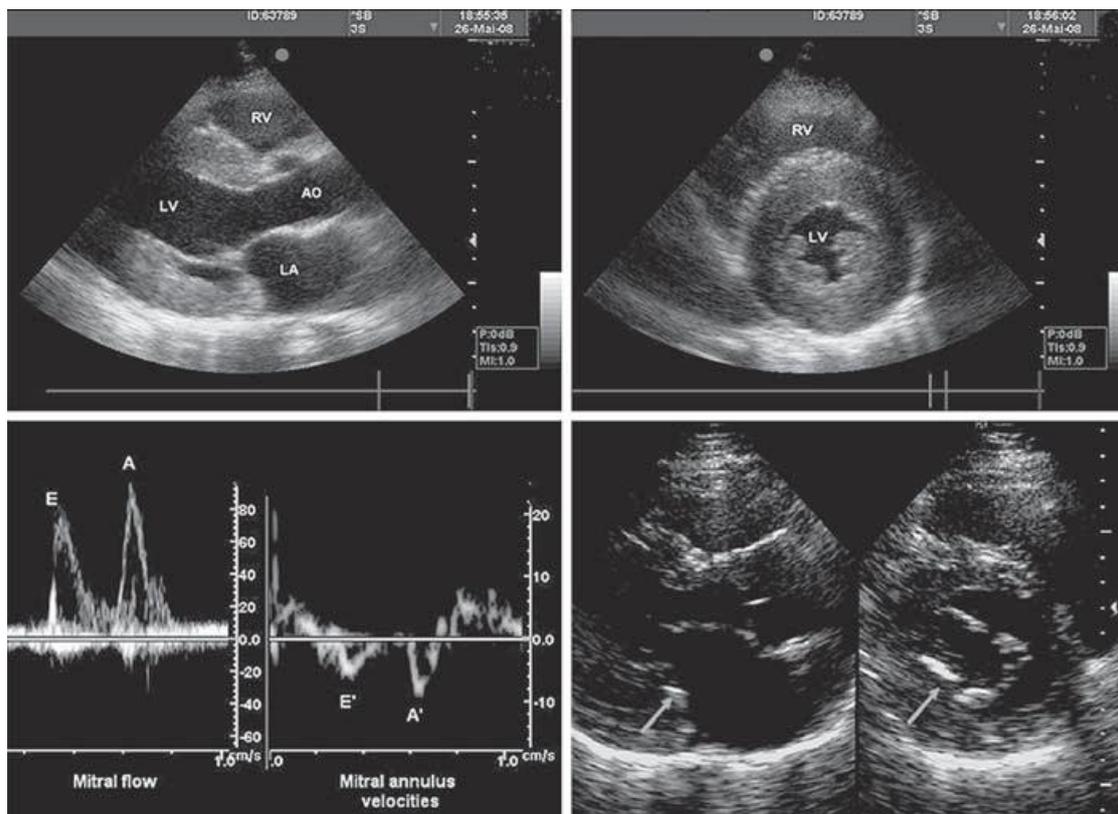


Fig-1: Representative images of echocardiographic findings commonly observed.

(a) Parasternal longitudinal view showing a pattern of concentric left ventricular hypertrophy. (b). Parasternal transversal view of the same patient. (c). Classic pattern of abnormal relaxation, pointing to a mild diastolic dysfunction (left: early (E) and late (A) diastolic mitral flow velocities showing inversion of the E/A ratio; right: early (E') and late (A') mitral annulus velocities confirming the abnormal relaxation). (d). Mitral annulus calcification. LV = Left ventricle; RV = right ventricle; AO = aorta; LA = left atrium.

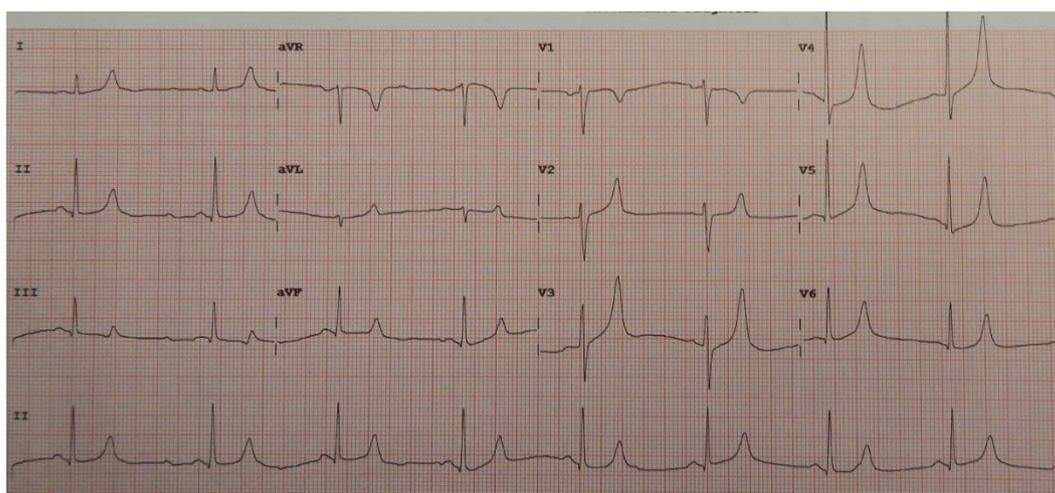


Fig-2: ECG changes of hyperkalemia in a patient with CKD

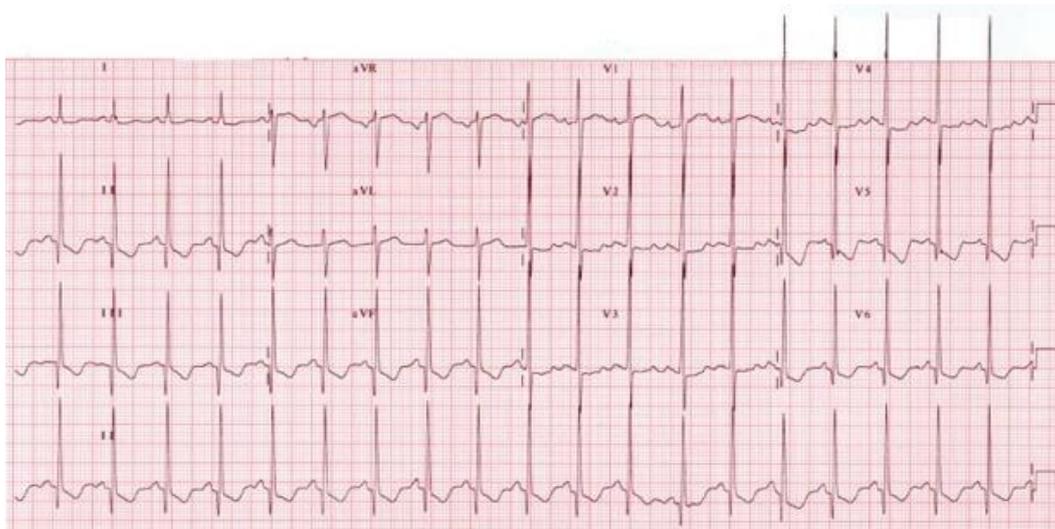


Fig-3: ECG showing left ventricular hypertrophy in a patient with CKD.

CONCLUSION

Electrocardiography and echocardiography are the non-invasive tools which can be used to identify cardiovascular disease early in the course of CKD. LVH was the most common echocardiographically detected abnormality. Diastolic dysfunction was present in 50% of cases and systolic dysfunction in 27.5% of cases. Periodic echocardiographic examination for diagnosis and treatment of cardiac abnormalities in patients with CKD is highly recommended.

REFERENCES

1. Heimdahl A, Støylen A, Torp H, Skjaerpe T. Real-time strain rate imaging of the left ventricle by ultrasound. *J Am Soc Echocardiogr* 1998;11:1013e9.
2. Go AS, Chertow GM, Fan D, McCulloch CE, Hsu CY. Chronic kidney disease and the risks of death, cardiovascular events and hospitalization. *N Engl J Med* 2004;351:1296e305.
3. Bargman JM, Skorecki K. Chronic Kidney disease. In: Longa DL, Fauci AS, Kasper DL, Hauser SL, Jameson JL, Loscal J, editors. *Harrison's principles of Internal medicine*, 18th ed New York: McGrawhill. 2012 p. 2308-21.
4. Prutkin JM. ECG tutorial: Basic principles of ECG analysis. UpToDate. 2013.
5. Salman shafi, Mahammad Saleem, Roshina Anjum, Wajid Abdullah Tahir Shafi. ECG abnormalities in patients with chronic kidney disease. *j Ayub Med Coll Abbottabad* 2017;29(1).
6. Agarwal S, Dangri P, Kalra OP, Rajpal S. Echocardiographic assessment of cardiac dysfunction in patients of chronic renal failure. *J Indian Acad Clin Med* 2003; 4:296-303.
7. Bullock RE, Amer HA, Simpson I, Ward MK, Hall RJ. Cardiac abnormalities and exercise tolerance in patients receiving renal replacement therapy. *Br Med J (Clin Res Ed)*. 1984 Dec 1;289(6457):1479-84.
8. Foley RN, Parfrey PS, Harnett JD, Kent GM, Martin CJ, Murray DC, Barre PE. Clinical and echocardiographic disease in patients starting end-stage renal disease therapy. *Kidney international*. 1995 Jan 1;47(1):186-92.
9. Singh NP, Nair M, Anuradha S, Kohli R, Agarwal SK. The cardiovascular and hemodynamic effects of erythropoietin in chronic renal failure. *The Journal of the Association of Physicians of India*. 2000 Mar;48(3):301-6.
10. Foley RN, Parfrey PS, Harnett JD, Kent GM, Murray DC, Barre PE. Impact of hypertension on cardiomyopathy, morbidity and mortality in end-stage renal disease. *Kidney international*. 1996 May 1;49(5):1379-85.
11. McGregor E, Jardine AG, Murray LS, Dargie HJ, Rodger RS, Junor BJ, McMillan MA, Briggs JD. Pre-operative echocardiographic abnormalities and adverse outcome following renal transplantation. *Nephrology, dialysis, transplantation: official publication of the European Dialysis and Transplant Association-European Renal Association*. 1998 Jun 1;13(6):1499-505.
12. P Dangri, S Agarwal, OP Kalra, S Rajpal. Echocardiographic assessment of left ventricular hypertrophy in patients of chronic renal failure. *Indian J Nephrol* 2003;13:92-97.
13. Levy D, Garrison RJ, Savage DD, Kannel WB, Castelli WP. Prognostic implications of echocardiographically determined left ventricular

- mass in the Framingham Heart Study. *N Engl J Med* 1990;27:1561-66.
14. Silberberg JS, Barre PE, Prichard SS, Sniderman AD. Impact of left ventricular hypertrophy on survival in end stage renal disease. *Kidney Int* 1989;36:286-90.
 15. Shapira OM1, Bar-Khayim Y. ECG changes and cardiac arrhythmias in chronic renal failure patients on hemodialysis. *J Electrocardiol* 1992 Oct;25(4):273-9.
 16. Ribeiro S, Ramos A, Brandao A, Rebelo JR, Guerra A, Resina C, Vila-Lobos A, Carvalho F, Remédio F, Ribeiro F. Cardiac valve calcification in haemodialysis patients: role of calcium-phosphate metabolism. *Nephrology, dialysis, transplantation: official publication of the European Dialysis and Transplant Association-European Renal Association*. 1998 Aug 1;13(8):2037-40.
 17. Maher ER, Young G, Smyth-Walsh B, Pugh S, Curtis JR. Aortic and mitral valve calcification in patients with end-stage renal disease. *The Lancet*. 1987 Oct 17;330(8564):875-7.
 18. Takenori Otsuka, Makoto Suzuki, Hisao Yoshikawa, Kaoru Sugi. Ventricular diastolic dysfunction in the early stage of chronic kidney disease. *Journal of Cardiology* 2009;54(2):199–204.
 19. Wizemann V, Kramer W, Schütterle G. The heart in end-stage renal failure: etiology, symptoms, and management of uremic heart disease. S Karger Pub; 1986.